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Upscaling of methane exchange in a boreal forest using soil chamber measurements and high-resolution LiDAR elevation data

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Abstract

Forest soils are generally considered to be net sinks of methane (CH₄), but CH₄ fluxes vary spatially depending on soil conditions. Measuring CH₄ exchange with chambers, which are commonly used for this purpose, might not result in representative fluxes at site scale. Appropriate methods for upscaling CH₄ fluxes from point measurements to site scale are therefore needed. At the boreal forest research site, Norunda, chamber measurements of soils and vegetation indicate that the site is a net sink of CH₄, while tower gradient measurements indicate that the site is a net source of CH₄. We investigated the discrepancy between chamber and tower gradient measurements by upscaling soil CH₄ exchange to a 100 ha area based on an empirical model derived from chamber measurements of CH₄ exchange and measurements of soil moisture, soil temperature and water table depth. A digital elevation model (DEM) derived from high-resolution airborne Light Detection And Ranging (LiDAR) data was used to generate gridded water table depth and soil moisture data of the study area as input data for the upscaling. Despite the simplistic approach, modeled fluxes were significantly correlated to four out of five chambers with $R > 0.68$. The upscaling resulted in a net soil sink of CH₄ of $-10\mu\text{mol m}^{-2}\text{h}^{-1}$, averaged over the entire study area and time period (June-September, 2010). Our findings suggests that additional contributions from CH₄ soil sources outside the upscaling study area and possibly CH₄ emissions from vegetation could explain the net emissions measured by tower gradient measurements.

Keywords: methane (CH₄) fluxes, water table depth, topographic wetness index, soil moisture, soil temperature, methane consumption

1. Introduction

The only well characterized biospheric sink for CH₄ is oxidation by methanotrophic bacteria in soil (Harriss et al., 1982). Globally, this soil CH₄ sink was estimated to range between 28 and 32 Tg CH₄ yr⁻¹, which amounts to around 5% of the destruction of CH₄ by OH radicals in the troposphere (Kirschke, 2013). Forest soils are generally considered to be net sinks of CH₄ with higher uptake rates than grassland and arable land (Boeckx et al., 1997; Dutaur and Verchot, 2007). However, CH₄ production by archaeans usually dominates in anaerobic forest soil environments such as waterlogged soils (Christiansen et al., 2012; Jungkunst et al., 2008; McNamara et al., 2006). CH₄ production also takes place in well-aerated soils at anaerobic micro sites (Fischer and Hedin, 2002; Kammann et al., 2009) and in deeper soil layers where anaerobic conditions occur (Kammann et al., 2001). Hence consumption and production can occur simultaneously at one location and soil conditions will determine the direction of the net flux. Vegetation might also contribute to the CH₄ exchange of a forest. Trees have been found to transport CH₄ originating from soil water and to release it through the stem or foliage (Terazawa et al. 2007; Gauci 2010). Aerobic formation of CH₄ in green plants has also been observed (Keppler et al. 2006; Vigano et al. 2008), although the mechanisms governing plant CH₄ release are still discussed (Bruhn et al., 2012) and there is little evidence of plant emissions of CH₄ from in-situ studies (Sundqvist et al., 2012). On the contrary, Sundqvist et al. (2012) found evidence for plant uptake of atmospheric CH₄ from measurements on spruce, pine, birch and rowan in a boreal forest.

Soil CH₄ flux rates also vary considerably both spatially and temporally (Christiansen et al., 2012; Ishizuka et al., 2009; Konda et al., 2008; Lessard et al., 1994; Reay et al., 2005; Yu et al., 2008). Spatial variability in soil CH₄ fluxes can be due to variability in soil moisture, soil

texture, and water table depths, factors that are dependent on topography, vegetation, and soil type, for example. Soil moisture (Castro et al., 1994; Guckland et al., 2009; Lessard et al., 1994) and soil texture (Dorr et al., 1993; Dutaur and Verchot, 2007; Ishizuka et al., 2009) alters soil diffusivity, which controls the rate at which atmospheric CH₄, and oxygen are supplied to the bacteria. Water table depth alters the relative extent of aerobic and anaerobic zones in soils. A rise of the water table leads to a decreased oxic soil zone and thus reduced CH₄ uptake (Kammann et al., 2001; Roulet et al., 1992). Changes in soil temperature and precipitation are also responsible for temporal variability in CH₄ exchange. Increases in temperature stimulate the activity of both methanogens (Yvon-Durocher et al., 2014) and methanotrophs (Crill et al., 1994; King and Adamsen, 1992), although methanogens benefit more (Dunfield et al., 1993). Other factors that have been found to influence soil CH₄ exchange in forests are soil pH (Weslien et al., 2009) and nitrogen availability (Steudler et al., 1989).

In situ chamber measurements and soil incubations in laboratories have long been the dominant methods for studying CH₄ exchange in forests, although larger scale micrometeorological methods are gaining in popularity (Nicoloni et al., 2013). While CH₄ exchange occurs and is often measured at the centimeter scale, it varies globally, and has a significant influence on biospheric-atmospheric interactions and feedbacks associated with climatic change (Schimel and Potter, 1995). Appropriate upscaling of CH₄ exchange from chamber-based point measurements will allow scientists to better understand the contribution of methane from soil and plant environments measured using eddy covariance/micrometeorological methods with extension to model estimates of regional to global CH₄ budgets (Hashimoto et al., 2011; Marushchak et al., 2013; Schimel and Potter, 1995). A few studies have upscaled CH₄ fluxes using simple extrapolations of chamber measurements or soil incubations from a few

locations multiplied by site area. However these methods do not consider the spatial heterogeneity of forest soil texture or type, or topographical variability, which may greatly influence wetting and drying regimes, and therefore CH₄ fluxes. Global and regional estimates of soil CH₄ sink strength use soil texture classes (Dorr et al., 1993; Dutaur and Verchot, 2007), land use type (Grunwald et al., 2012), ecosystem class and/or climatic zones (Dutaur and Verchot, 2007) to spatially parameterize CH₄ exchanges. However, regional models often fail to incorporate the spatial heterogeneity within each class, including fuzzy boundaries between classes. This results in inaccurate characterization of classes, and especially within the sometimes broad transition zones between classes (Matson et al., 1989). These issues may be overcome by incorporating process-based models of CH₄ consumption driven by gaseous diffusion or diffusion in combination with microbial activity (Curry, 2007; Del Grosso et al., 2000; Ridgwell et al., 1999). Some process-based models do not account for production of CH₄ and are not applicable to soils that seasonally shift from net sinks to net sources (Del Grosso et al., 2000). Process-based models can become exceedingly complex, requiring detailed inputs of spatio-temporally varying climate, vegetation and soil physiochemical properties (Hashimoto et al., 2011). More simple, empirical models have been developed for site-specific applications. Castro et al (1994) found that soil moisture, as the only explanatory variable, could satisfactorily predict CH₄ fluxes at locations within a temperate forest. Christiansen et al. (2012) used spatial variability in soil moisture and water table depths derived from elevation data to upscale CH₄ fluxes from manual chamber measurements to site scale at two temperate deciduous forests.

At the Norunda boreal forest site in central Sweden, chamber measurements of soils and vegetation indicate that the site is a net sink of CH₄ (Sundqvist et al., 2012, 2014b), while

gradient measurements above the forest canopy indicate that the site is a net source of CH₄ (Sundqvist et al., 2015). The aim of this study was to quantify soil CH₄ exchange for the entire site (100 ha) by upscaling soil CH₄ exchange through developing an empirical model for a mature coniferous forest based on automated chamber observations with a high temporal resolution, in combination with high-resolution LiDAR elevation data. The model will also serve as a mean to further examine the discrepancy between results obtained from chamber measurements and tower gradient measurements. In correspondence to findings of Christiansen et al. (2012), Fiedler et al. (2005) and Grunwald et al. (2012), we hypothesize that emissions from wet patches scattered at the site may exceed the uptake in well-aerated parts of the soil and hence even relatively small source areas may shift a larger area from a sink to a source (Fiedler et al., 2005).

2. Method

2.1 Site description

Upscaling of soil CH₄ exchange was completed for a 100 ha area at the Norunda site, 60°5'N, 17°29'E, in central Sweden from July through September 2010 during coincident chamber and tower gradient measurements. The Norunda site is situated at the southern edge of the boreal forest zone and is comprised of 120 years old mixed pine (*Pinus sylvestris*) and spruce (*Picea abies*) trees. The forest was thinned in 2008 within the NE to SW sectors surrounding the measurement tower to a radius of 200 m, which decreased the leaf area index within this area from 4.8 to 2.8 m²m⁻². Trees within the SW to NE sectors have not been thinned nor fertilized in the last few decades. Soil are comprised of glacial till, classified as dystic regosol (Lundin et al., 1999) and include an organic layer of about 3-10 cm. The area within 500 m radius of the measurement tower is relatively flat, with elevation ranges from 40-52 m above sea level (Figure 1). Since 1843, the water table in the area has been artificially lowered as a result of several ditches surrounding the forest. The last known ditch installation was in 1980. Mean air temperature measured at Uppsala climate station, 30 km south of Norunda, was 6.5° C and mean precipitation was 576 mm (1980-2010).

2.2 Instrumentation

In this study, eight CH₄ chambers are used: three chambers (T1-T3) were located in the thinned section, and five (U1-U5) were located in the undisturbed section of the forest. In areas of higher water table, CH₄ exchanges were measured, using a single 'floating' chamber positioned on standing shallow water in the thinned section of the forest.

Due to equipment limitations, measurements were conducted at one section at a time and were averaged from hourly chamber measurements to daily periods from 1 August to 30 September 2009 at the thinned plot, and 7 July to 30 September 2010 at the undisturbed plot. The soil chambers each had a volume of 110 l and covered a surface-area of 0.2 m². Concentrations in the chambers were measured for 6 minutes after chamber closure with a high precision off-axis integrated cavity output spectroscopy laser gas-analyser (DLT-100, Los Gatos Research). A small fan was installed in each chamber to ensure proper headspace mixing during closure. Air was circulated between the chamber and a manifold at a flow rate of 8-10 l min⁻¹. A sub-sample of the air stream was passed through the gas analyzer at a flow rate of 1.2 l min⁻¹. The rate of change in CH₄ concentration was determined through a linear fit to the first 2 minutes of concentration data and averaged over hourly periods. CH₄ fluxes were calculated as

$$F_{CH_4} = \frac{dC}{dt} \frac{V}{A}, \text{ where } C \text{ (}\mu\text{mol m}^{-3}\text{), is molar density, } V \text{ (m}^3\text{) is chamber volume and } A \text{ (m}^2\text{) is}$$

the ground surface area covered by the chamber. Only measured fluxes were used for upscaling, i.e. CH₄ flux data was not gap-filled. For further details on the chamber measurements, see Sundqvist et al. (2014b). Soil temperature was measured continuously at a depth of 5 cm below the soil surface within the chambers using type-T thermocouples. Soil moisture was also measured continuously and averaged within the top 5 cm of the soil within chambers using Ml-2x ThetaProbe from DeltaT Devices and at additional positions throughout the study area.

2.3 Empirical model for upscaling of CH₄ exchange

Upscaling of soil CH₄ exchanges was derived using a multiple linear regression analyses (Draper and Smith, 1998) based on typical control variables: soil temperature, soil moisture and water table depth as they compare with hourly average chamber measurements. The resulting

regression model was assumed to be valid as long as the water table was greater than 5 cm below the soil surface such that an oxic soil zone could enable CH₄ consumption by methanotrophs. For higher water tables a median value of 40 $\mu\text{mol m}^{-2}\text{h}^{-1}$ obtained from the floating chamber measurements was used. The mean value of the floating chamber was as large as 120 $\mu\text{mol m}^{-2}\text{h}^{-1}$ due to several ($>1000 \mu\text{mol m}^{-2}\text{h}^{-1}$) flux values, possibly related to ebullition. Different combinations of chamber measurement were tested when the empirical model was derived through regression analyses. Three chambers were always used for model development while the other five were used for model evaluation. The model that showed the best fit between input data and regression curve and best predicted fluxes was selected for upscaling. Sensitivity analyses of model parameters were made by local sensitivity analyses by varying one variable at the time and keeping the other fixed (Hamby, 1994). Each parameter was changed by 20% at the time and the change in the model output was quantified.

2.4 Model input

For upscaling of soil CH₄ exchange to site level, spatial information on soil moisture, water table depth and soil temperature were needed as input. Daily average soil temperatures at the site were relatively constant in space, hence only temporal variations in temperature were accounted (Table 1). Spatial information of soil moisture and water table depth were estimated using airborne-LiDAR data collected in June 2011.

LiDAR point cloud data were first classified to ground and non-ground returns using TerraScan (TerraSolid, Inc.). Ground-classified returns were rasterized using an inverse distance weighted algorithm (Hopkinson et al., 2005) to generate a digital elevation model (DEM) with a cell resolution of 1 m (Figure 1). A topographic wetness index (TWI) was derived

from the DEM (Beven and Kirkby, 1979), and used to estimate unsaturated zone soil moisture according to Lin et al., (2006), Schmidt and Persson, (2003), and Western et al., (1999). Further, Beldring et al., (1999) demonstrated that within some forested areas, the spatial patterns of soil moisture can be estimated more accurately using topography than with other hydrological state measurements, as the latter are often too variable when used with sparsely distributed measurements. TWI is calculated as $TWI = \ln(A / \tan \beta)$, where A is the upslope area (m²) being drained through the position of interest, and β is the slope angle (degrees) of that position (Beven and Kirkby, 1979). In the present study, the flow routing and the drainage area (A) have been calculated in a multiple flow direction algorithm that takes the shape of the slope into account (Pilesjö et al., 1998; 2006). TWI was converted to soil moisture based on the linear regression between TWI and soil moisture measured at 12 site locations in 2010 (Figure 2). TWI was also compared with soil moisture measured in 2013 as a separate evaluation of TWI as a proxy for soil moisture at the site. Soil moisture data in 2013 were sampled at several positions as oppose to other years (Figure 2).

The DEM was also used to estimate the spatial variability of water table depths based on the assumption that topographically low areas are likely to have water table depths nearer to the surface than topographically higher positions (e.g. Lamb et al., 2001, Molénat et al., 2005, Seibert et al., 1997).

Since continuous measurements of water table depth were available only at a single location in the study area a smoothed ground surface was used to represent the water table. The smoothened ground surface area was lowered in each cell with the water table depth obtained from the one measurement location. The elevation values' spatial autocorrelation were plotted in semi-variograms in which the spatial dependency of elevation was estimated. The geomorphology of

the area was studied in aerial imagery and within the DEM. The average measure from valley bottom to hilltop to valley bottom of the ground surface and the results of the spatial auto-correlation formed the base of the smoothing process. The smoothened ground surface was obtained by applying a low-pass filter that averaged the elevation values up to a distance where the spatial auto-correlation of elevation declined.

Validation of the spatial distribution of the depth to water table and soil moisture was determined using LiDAR-derived vegetation heights as a proxy indicator, assuming that trees are often not as tall in wet soils compared to drier soils (Heiskanen and Makitalo, 2002; Koslowski, 1997; Polacek et al., 2006). Spatial variations in tree heights were determined using a digital surface model (DSM) derived from LiDAR data. The DSM was determined by gridding the maximum height of classified non-ground laser pulse returns within a 3 meter search radius and subtracted from the DEM to determine vegetation height relative to ground surface. The comparison between tree height and depth to water table and soil moisture could be made for an area within 250 m radius from the tower where the trees are the same age.

2.5 Gradient measurements

Gradients were measured between 31.7m and 58.5 m, and between 31.7 m and 101.6 m at the main tower at Norunda with a laser gas analyser (DLT-100, Los Gatos Research). Here we used the gradient measured between 31.7 m and 58.5 m since footprint analyses using the model by Kljun et al (2004) showed that these levels were more representative for the upscaled area than the gradient including the highest level (Sundqvist et al., 2015). Tower CH₄ fluxes were calculated in two different ways; the combined eddy covariance and gradient method (ECG) and the modified Bowen-ratio method (BR)

(Sundqvist et al., 2015). In the ECG method the CH₄ flux is calculated according to Ficks law of diffusion:

$$F_{CH_4}(ECG) = -\frac{1}{V} K_c \frac{\Delta C}{\Delta z} \quad (1)$$

where V (m³ mol⁻¹) is the volume of one mole of air and C is CH₄ concentration (μmol mol⁻¹).

The turbulent diffusivity K_c is derived from the Monin Obukhov similarity theory,

$$K_c = \frac{ku_*\Delta z}{\ln \frac{z_2}{z_1} - \psi_t\left(\frac{z_2}{L}\right) + \psi_t\left(\frac{z_1}{L}\right) - \int_z^{z_*} \Phi_t(1 - \phi_t) \frac{dz}{z}} \quad (2)$$

where $k = 0.4$ is von Karman's constant, u_* (m s⁻¹) is friction velocity, z_2 (m) is upper air intake height, z_1 (m) is lower air intake height, Δz is the difference between the intake heights, L is the Obukhov length and ψ_t is the diabatic correction function for heat profiles according to Högström (1988). u_* and L were obtained from the eddy covariance measurements. The function Φ_t denotes the diabatic correction function for gradients. Φ_t is integrated from the height z inside the roughness sublayer to the top of the roughness sublayer ($z_* = 35.9$ m). The roughness sublayer effects are expressed through ϕ_t (Mölder et al., 1999; Physick and Garratt, 1995). All heights take the displacement height into account.

The modified Bowen ratio method is based on the assumption of identical diffusivities for CO₂ and CH₄ fluxes. If the CO₂ flux (F_{CO_2}) is known, e.g., the CH₄ exchange F_{CH_4} can be derived (Moncrieff et al., 1997) as

$$F_{CH_4}(BR) = F_{CO_2} \frac{\Delta C_{CH_4}}{\Delta C_{CO_2}} \quad (3)$$

262 A measurement uncertainty was calculated for each half hourly value based on estimated
263 uncertainty in the ingoing variables. Assuming independence between the errors, the error in the
264 mean value for the investigated period was propagated according to the simplified method (Hu,
265 1966) for both the ECG and BR methods. The statistical uncertainty was calculated as the
266 standard error of the mean. The total error in the mean was estimated as the sum of these
267 uncertainties.

3. Results

3.1 Empirical model

The combination of chamber data used for developing the model had a clear impact on the goodness of fit of the empirical model (Table 2). This was also the case for the correlation between modelled fluxes and fluxes from chambers used for evaluation of the model. The model with best fit (R^2) to input data and highest correlation with evaluation data (R) was chosen for upscaling of the CH_4 exchange. The model with the overall best performance had an R^2 of 0.8 and R values of 0.85, 0.72, 0.94, 0.68 and 0.15 (Table 2) for U1, U2, U5, T1 and T3, respectively. All R values except the one for T3 were significant. The model equation for estimating the net CH_4 exchange ($\text{CH}_{4\text{exchange}}$) was $\text{CH}_{4\text{exchange}} = -0.32T + 3.5W + 0.08SM - I$, where T is soil temperature, W is water table depth, SM is soil moisture and I is the regression intercept which equaled $2.8 \mu\text{mol m}^{-2} \text{h}^{-1}$. Uptake values ($\mu\text{mol m}^{-2} \text{h}^{-1}$) were overestimated for U1, U5 and T3 and underestimated for U2 (Figure 3). Sensitivity analyses showed that the model was equally sensitive for changes in water table depth and soil temperature and less sensitive to changes in soil moisture. A 20 % change in parameter values resulted in 9.1%, 9.2% and 2.7 % change in net CH_4 exchange for water table depth, soil temperature and soil moisture, respectively.

3.2 TWI and soil moisture

TWI and soil moisture were positively correlated with R values ranging from 0.1-0.7 in 2013 and 0.2-0.7 in 2010 (Figures 4a-b). The correlation coefficients were higher on average in 2013 (Figure 4a). Correlations were not significant due to the limited number of soil moisture

measurements. P-values for 2010 were generally < 0.2 (Figure 4c). When soil moisture decreased after precipitation, the correlation between TWI and soil moisture increased slightly (Figure 4d). Spatio-temporal variability in soil moisture generated from the relationship between TWI and soil moisture measurements was on average between 14.7- 22.3 % for 96 % of the data (Figure 5a). Daily input data from soil moisture measurements varied between 5-35 %. A map of tree heights in the study area is shown in Figure 5b. It can be seen that areas with lower than average tree heights coincide with areas with higher than average soil moisture in the area within 250 m radius from the tower where trees are the same age. There was a significant correlation between soil moisture and tree height with a correlation coefficient of -0.16.

3.3 Water table depth

For each 1 m by 1 m cell, the spatial dependency of elevation on the water table was estimated to be 80 m. A water table surface was generated applying a low-pass filter on the DEM that averaged the elevation values taking 80 m surroundings into account. The ground water measurements were used to let this filtered water table fluctuate temporarily. Spatio-temporal variability in water table ranged on average from -5 m below ground to 0.7 m above ground if excluding outliers, with 2 and 98 percentiles of -2.9 m and -0.27 m. A few wet patches with standing water are shown in the map of modelled water table depth averaged over the study time period (Figure 5c). In figure 5b it can be seen that in areas within 250 m from the tower, lower than average tree heights coincide with areas with higher than average water tables. There was a significant correlation between depth to water table and tree height with a correlation coefficient of -0.29.

3.4 Upscaled CH_4 exchange

Averaged modelled net CH_4 exchange (derived as described in Section 2.2) of the entire study area and time period is $-10.0 \mu\text{mol m}^{-2} \text{h}^{-1}$ with a standard deviation of $0.8 \mu\text{mol m}^{-2} \text{h}^{-1}$.

However, the average CH_4 exchange for the entire measurement period varies spatially between uptake values of $-21 \mu\text{mol m}^{-2} \text{h}^{-1}$ and the assigned production value of $40 \mu\text{mol m}^{-2} \text{h}^{-1}$ for areas with standing water (Figure 5d). Net average emissions are found for 1.75% of the study area.

Time series of daily averaged modelled CH_4 exchange, soil moisture, soil temperature and water table depth are shown in Figure 6. Over the course of the measurement period, net CH_4 uptake decreased along with decreased soil temperatures.

4. Discussion

4.1 Evaluation of model and upscaling

Although the empirical model for upscaling of CH₄ exchange only includes data from three chamber locations and is driven by the three variables, soil temperature, depth to water table and soil moisture, this method shows good agreement between modelled and measured fluxes, in general. However the model is unsuccessful in capturing local deviations in CH₄ flux behavior such as the peak emission followed by a shift from source to sink of chamber T3 (Figure 3). The peak in CH₄ emissions may be due to high soil moisture level following precipitation (Sundqvist et al., 2014b), with the shift to a sink when the soil dries up.

Since the model is less sensitive to changes in soil moisture than to changes in water table depth and soil temperature (cf. Eq. 1), soil moisture driven events might not be fully captured.

Modelled estimates of soil moisture and water table depths derived from LiDAR data agrees well with the qualitative picture we have of the study area. Moreover, the shorter trees are found in areas with higher soil moisture and near surface water table, which support the hydrological patterns generated from the DEM (Heiskanen and Makitalo, 2002; Koslowski, 1997; Polacek et al., 2006). Future studies should include more measurements of soil moisture and water table depths covering a larger area for a more quantitative evaluation. It would also be desirable to evaluate the upscaling by measuring CH₄ fluxes with soil chambers at several locations in the area.

4.2 Upscaled CH₄ exchange

The results of the modelled/upscaled soil CH₄ exchange indicates that this forest is an overall sink of -10.0 μmol m⁻²h⁻¹. Even if the assigned production areas of 40 μmol m⁻²h⁻¹ (for water

table depths higher than 5 cm below the soil surface) were extended to all water table depths higher than 30 cm below the soil surface, the area remained a net sink of $-9.1 \mu\text{mol m}^{-2}\text{h}^{-1}$. According to the chamber measurements, the average CH_4 exchange for the undisturbed forest and the thinned forest were $-10 \mu\text{mol m}^{-2} \text{h}^{-1}$ and $-5 \mu\text{mol m}^{-2} \text{h}^{-1}$, respectively (Sundqvist et al., 2014b), and hence agreed well with the modelled values. Results from the modelled site-wide water table depth show that 90 % of the area had a water table lower than -0.7 m, which should be well below the critical levels for net CH_4 emissions. Christiansen et al. (2012) found that maintaining a ground water table depth below 0.5 m throughout their study area would shift their site from a source of CH_4 to a sink of CH_4 . Assuming that an uptake of $-10 \mu\text{mol m}^{-2}\text{h}^{-1}$ is reasonable in well aerated soils, a net production area of more than 20 % of the total area, with average fluxes of $40 \mu\text{mol m}^{-2}\text{h}^{-1}$, would be necessary to shift the site from a sink to a source. The size of this area would of course decrease if the CH_4 production rates approached levels of wetland fluxes. In general, uptake rates decreased from July to September (Figure 6) and it is possible that additional data from wetter and colder seasons would alter the findings.

4.3 Water table depth

We used filtered elevation data derived from LiDAR measurements to derive the general pattern of the water table in the study area and regulated the absolute depths of the modelled water table with measurements from one single location. Other studies have modelled water tables from TWI, which may be related to topography, depending on soil type (Buttle et al. 2004). Siebert et al. (1997) concluded that one single location of observation of water table depth was sufficient for the calibration of an index to predict groundwater spatially distributed in a spruce forested catchment in the West of Sweden. However, we are aware of an area with standing water in

summer 2010, which was not accurately captured by the model, although the generated water table map and soil moisture map show higher than average values for this area. The generation of the water table depth is based solely on the DEM. In nature, water table depth is also dependent on soil properties such as large pore structure, impermeable clay layers and sub-surface lateral flow in hillslopes (Grip and Rodhe, 1994; Ward and Trimble, 2004). Clay layers in Norunda soils have been observed at a few locations, but their spatial extent and general depth are unknown.

Moreover, the water table map was developed under conditions where the observed water table depth ranged from 0.9 m-1.7m below soil surface. Jungkunst et al. (2008) found that CH₄ emissions of a hydromorphic forest soil increased drastically when the water table was 10-20 cm from the soil surface. Fiedler and Sommer (2000) determined a threshold depth of 15 cm of the water table for wetland soils; above this threshold, production dominated the CH₄ exchange. Since the control of CH₄ production at high water tables differs from drivers of CH₄ consumption and mainly depends on substrate availability and soil temperature (Christensen et al., 2003), an additional model developed for high water tables might improve the upscaling.

4.4 Soil moisture

Observed soil moisture was well represented by the model with variations between 5% and 35%. However, the average spatial variation of soil moisture generated from TWI was lower. The porosity of the soil in Norunda is around 40% and in areas with standing water, soil moisture should reach the value of the porosity. Only 0.5 % of the cells had average soil moisture above 24% and no cells had average values above 31%. These low values may point to the fact that wet areas with observed standing water were situated in parts of the landscape which are represented

as flat or convex in the DEM. The water routing algorithm creates a pattern, which is based only on the topography. The model might then disperse water over larger areas rather than concentrating it due to low differences in elevation values. In reality, however, micro-topography and soil properties may drive the water distribution.

4.5 Upscaling soil CH₄ exchange versus gradient measurements

The results from upscaling of soil CH₄ exchange show a net consumption of about -10 $\mu\text{mol m}^{-2}\text{h}^{-1}$ while the tower gradient measurement indicates the site is a net CH₄ source of about 3.4 $\mu\text{mol m}^{-2}\text{h}^{-1}$ for the same time period (Sundqvist et al., 2015). The ECG method resulted in a mean emission of $5.2 \pm 0.64 \mu\text{mol m}^{-2}\text{h}^{-1}$ while the BR method gave $1.6 \pm 0.73 \mu\text{mol m}^{-2}\text{h}^{-1}$. The errors of these two methods does not overlap but both indicate that the forest is a source of methane. The difference between the methods is most likely caused by uncertainty in the methods itself. Here are many empirical corrections involved especially in the ECG method that might be the cause of this discrepancy.

Unfortunately the uncertainty in the upscaled soil emissions cannot be easily done since we do not have enough spatial data on e.g. soil properties available. The rough sensitivity analyses made in section 4.2 is however a strong indication that the soil is a sink of methane and not a source.

The study area included the outskirts of a clear-cut (Figure 2). The water table in this part of the study area was increased due to decreased evapotranspiration following clear-cutting. The model did not capture this local effect since water table depth was calculated from elevation data and a single measurement location within the forest. Chamber measurements on the clear-cut from October -November 2010 showed average emissions of 15 $\mu\text{mol m}^{-2}\text{h}^{-1}$ (Sundqvist et al.,

2015). Emissions from the clear-cut may be part of the reason for the discrepancy between the upscaled soil chamber data and the gradient measurements. It is also possible that emissions from the part of the clear-cut located outside of the upscaled boundary and other sources outside the study area contribute to the tower gradient measurements.

The gradient measurements represent the net exchange of CH₄ above the canopy and hence it is not possible to distinguish fluxes at soil level from an exchange of CH₄ coupled to vegetation. CH₄ emissions by vegetation (Terazawa et al. 2007; Gauci 2010; Keppler et al. 2006; Vigano et al. 2008) could partly explain the discrepancy between upscaled chamber data and gradient measurements, if these contribute significantly to the net CH₄ exchange at Norunda. However, so far only a net uptake of CH₄ by vegetation has been measured at a few locations at Norunda (Sundqvist et al. 2012).

5 Conclusions

A simple empirical model for upscaling of soil CH₄ exchange at a boreal forest site was developed from chamber measurements, driven by soil temperature, water table depth and soil moisture. High resolution LiDAR elevation data formed the basis for the mapping of water table depth and soil moisture of the study area, while vegetation heights were used as a proxy indicator for spatial variations of average soil moisture regime. A significant correlation was found between the topographical wetness index (TWI) derived from the LiDAR data and soil moisture measured at the study area, which enabled a conversion of TWI to soil moisture. Modelled CH₄ fluxes were in good agreement with measured fluxes at four out of five chamber locations used for evaluation. An improvement of the model could be achieved by a better representation of high water tables. The results from upscaling of CH₄ exchange show a net uptake of CH₄ of -10

448 $\mu\text{mol m}^{-2}\text{h}^{-1}$. This is in contrast to gradient measurements above the canopy in the center of the
449 study area, which yielded net CH_4 emissions for the same time period (Sundqvist et al., 2015).
450 Emissions were only found from 1.75 % of the surface of the study area. This suggests that the
451 net emissions measured by the tower gradient method above the tree canopy probably originated
452 from sources outside the study area and possibly also from CH_4 emissions by the vegetation.
453

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Tables

Table 1.

Soil temperature variability at Norunda. Here *Mean* and *Std* are the mean and standard deviation of the soil temperature respectively at *n* number of measurement locations within the study area. The measurements were taken in 2014.

	Mean (°C)	Std (°C)	n
July (undisturbed forest)	17.2	2.1	21
July (thinned forest)	17.9	1.4	12
September (undisturbed forest)	8.3	1.0	21
September (thinned forest)	7.8	0.9	12

Table 2

Evaluation of empirical models for upscaling of CH₄ exchange.

ID is the combination of chambers used for model development. R² is the adjusted coefficient of determination for the multiple linear regression analyses between model and chamber input data.

B(T), B(W) and B(SM) are the coefficients for soil temperature, water table depth and soil

moisture, respectively, obtained from the multiple linear regression analyses. I is the regression

intercept. R is the Pearson correlation coefficient for modelled CH₄ exchange and chamber data.

*) Not significant at p > 0.05. n.i) Not included due to non-significance. -) Chamber was used for

model development and is therefore not evaluated.

ID	R ²	B (T)	B (W)	B (SM)	I	R (U1)	R (U2)	R (U3)	R (U4)	R (U5)	R (T1)	R (T2)	R (T3)
U4, U5, T1	0.68	-0.43	5.8	n.i	4.8	0.79	0.79	0.81	-	-	-	0.71	-0.33*
U4, U3, T1	0.80	-0.31	3.1	0.08	3.7	0.80	0.78	-	-	0.96	-	0.63	-0.43*
U4, U2, T1	0.58	-0.30	1.9	0.09	-6.3	0.85	-	0.87	-	0.94	-	0.48	0.13
U4, U2, T1	0.55	-0.42	7.2	n.i	7.2	-	0.78	0.81	-	0.94	-	0.71	-0.21*
U4, U5, T2	0.71	-0.20	2.2	0.03	-4.6	0.84	0.74	0.88	-	-	0.62	-	-0.09*
U4, U3, T2	0.80	-0.32	3.5	0.08	-2.8	0.85	0.72	-	-	0.94	0.68	-	0.15*
U4, U2, T2	0.57	-0.33	3.6	0.05	-2.4	0.84	-	0.88	-	0.96	0.62	-	-0.09*
U4, U1, T2	0.54	-0.46	6.0	n.i	5.8	-	0.79	0.81	-	0.95	0.41	-	-0.34*
U4, U5, T3	0.56	n.i	8.1	0.16	0.60	0.69	0.61	0.74	-	-	0.08	-0.03*	-
U4, U3, T3	0.54	n.i	6.8	0.22	-4.6	0.71	0.58	-	-	0.67	0.58	-0.04*	-
U4, U2, T3	0.52	0.40	4.5	0.30	-15.3	0.43	-	0.36	-	0.18*	0.47	-0.23*	-
U4, U1, T3	0.53	n.i	8.9	0.15	2.3	-	0.62	0.74	-	0.68	0.54	-0.02*	-
U5, U3, T1	0.51	-0.37	n.i	0.05	-5.9	0.81	0.66	-	0.83	-	-	0.52	-0.46*
U5, U2, T1	0.28	-0.39	n.i	0.03	-5.7	0.80	-	0.72	0.80	-	-	0.52	-0.53*
U5, U1, T1	0.58	-0.49	4.7	-0.03	5.7	-	0.79	0.73	0.82	-	-	0.62	-0.53*
U5, U3, T2	0.53	-0.39	n.i	0.06	-5.7	0.81	0.65	-	0.83	-	0.41	-	-0.42*
U5, U2, T2	0.30	-0.42	1.3	n.i	-2.7	0.78	-	0.74	0.80	-	0.19	-	-0.57*
U5, U1, T2	0.59	-0.51	2.6	n.i	2.2	-	0.75	0.77	0.83	-	0.24	-	-0.54*
U5, U3, T3	0.44	n.i	n.i	0.32	-15.1	0.64	0.44	-	0.52	-	0.59	-0.06*	-
U5, U2, T3	0.39	n.i	n.i	0.33	-16.1	0.64	-	0.51	0.52	-	0.59	-0.06*	-
U5, U1, T3	0.48	n.i	3.4	0.24	-6.9	-	0.53	0.62	0.65	-	0.60	-0.05*	-
U3, U2, T1	0.50	-0.17	2.9	0.08	6.5	0.83	-	-	0.90	0.89	-	0.31	0.53
U3, U1, T1	0.25	-0.40	-1.6	0.03	-6.7	-	0.57	-	0.67	0.80	-	0.46	-0.56*
U3, U2, T2	0.49	-0.19	4.6	0.05	-3.1	0.80	-	-	0.93	0.89	0.66	-	0.37
U3, U1, T2	0.29	-0.42	-1.8	0.05	-7.0	-	0.57	-	0.70	0.81	0.26	-	0.53
U3, U2, T3	0.53	0.35	12.0	0.21	-3.4	-0.43*	-	-	-0.49*	-0.78*	0.25	-0.39*	-
U3, U1, T3	0.37	n.i	n.i	0.30	-14.2	-	0.44	-	0.52	0.51	0.59	-0.06*	-
U2, U1, T1	0.16h	-0.44	n.i	n.i	-3.8	-	-	0.68	0.72	0.87	-	0.50	-0.59*
U2, U1, T2	0.19	-0.46	n.i	n.i	-3.5	-	-	0.68	0.72	0.87	0.13	-	-0.59*
U2, U1, T3	0.35	n.i	5.7	0.20	-4.8	-	-	0.69	0.73	0.66	0.59	-0.04*	-

Figures

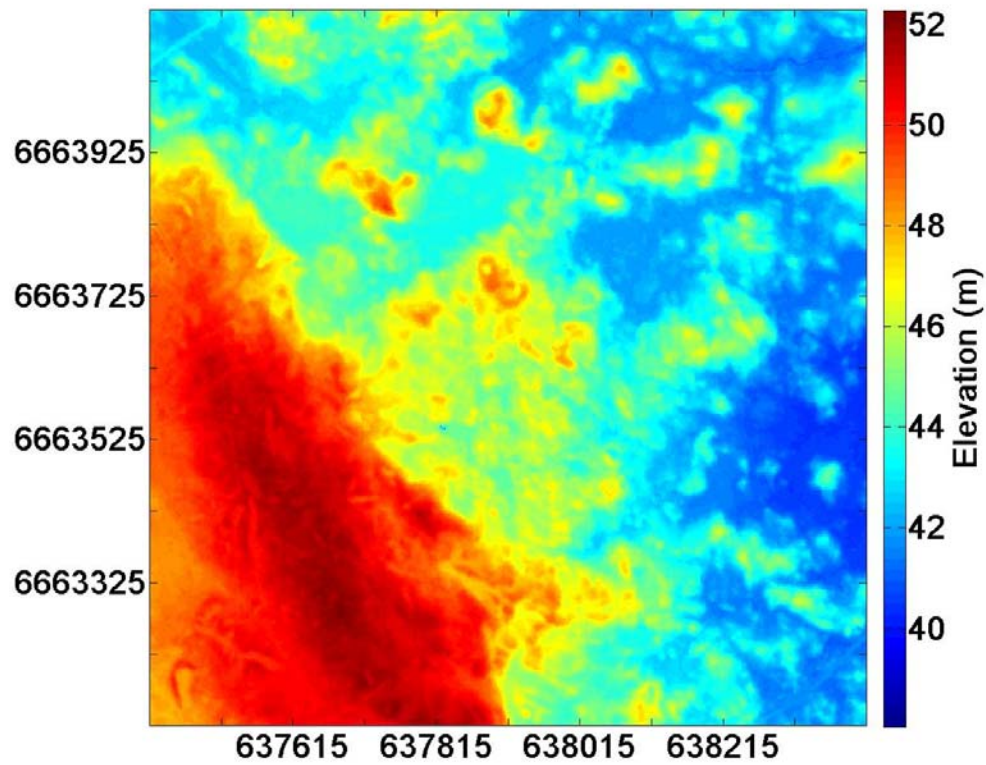


Figure 1. Digital Elevation Model (DEM) of the study area. Coordinates are given in UTM (WGS 84).

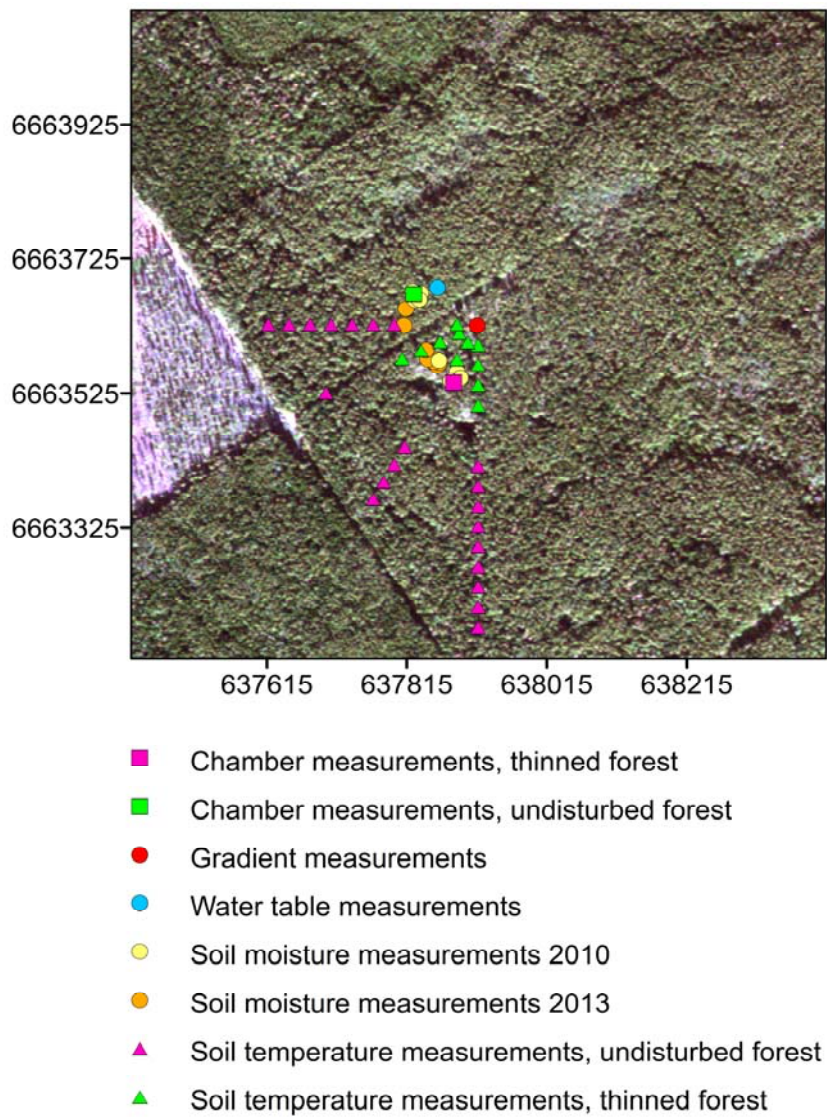


Figure 2. Photo of the study area showing locations for chamber measurements, soil moisture measurements, water table measurements and gradient measurements. The clear-cut can be seen to the left.

(Source: Geoeye-1© GeoEye Inc.<2009> Distributed by e-GEOS)

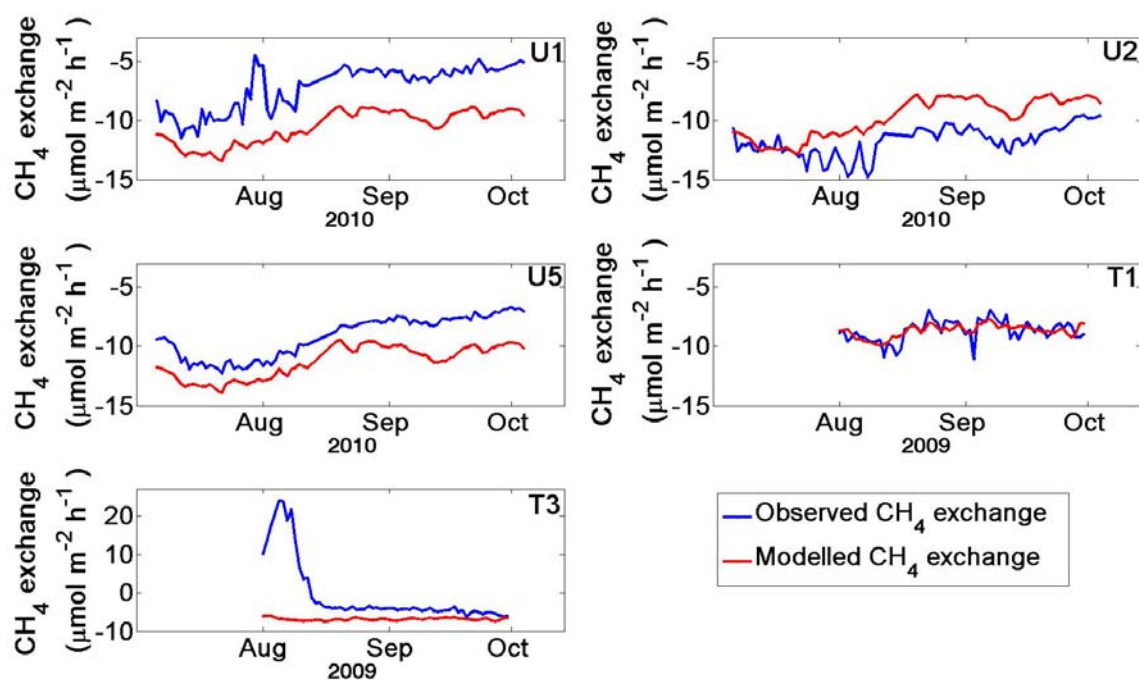


Figure 3. Comparison between modelled and observed CH₄ exchange for the evaluation chambers. Ux are chambers in the undisturbed forest and Tx are chambers in the thinned forest.

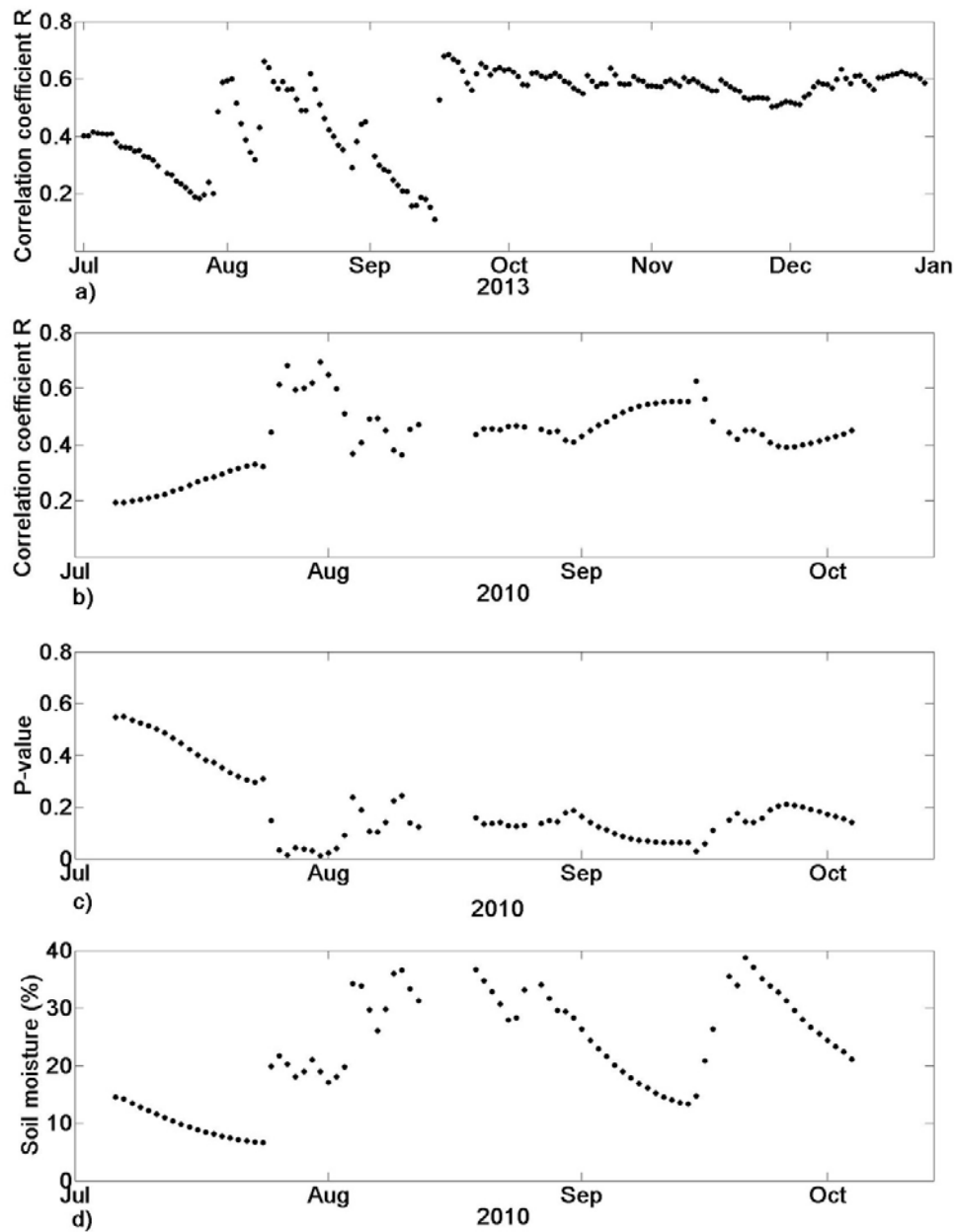


Figure 4. Average daily Pearson correlation between TWI and soil moisture measured at a) 9 locations July- December 2013; and b) 12 locations July-October 2010; Daily significance level for correlations between c) TWI and soil moisture measured at 12 locations July-October 2010; and d) daily soil moisture at one measurement location for the time period July-October 2010.

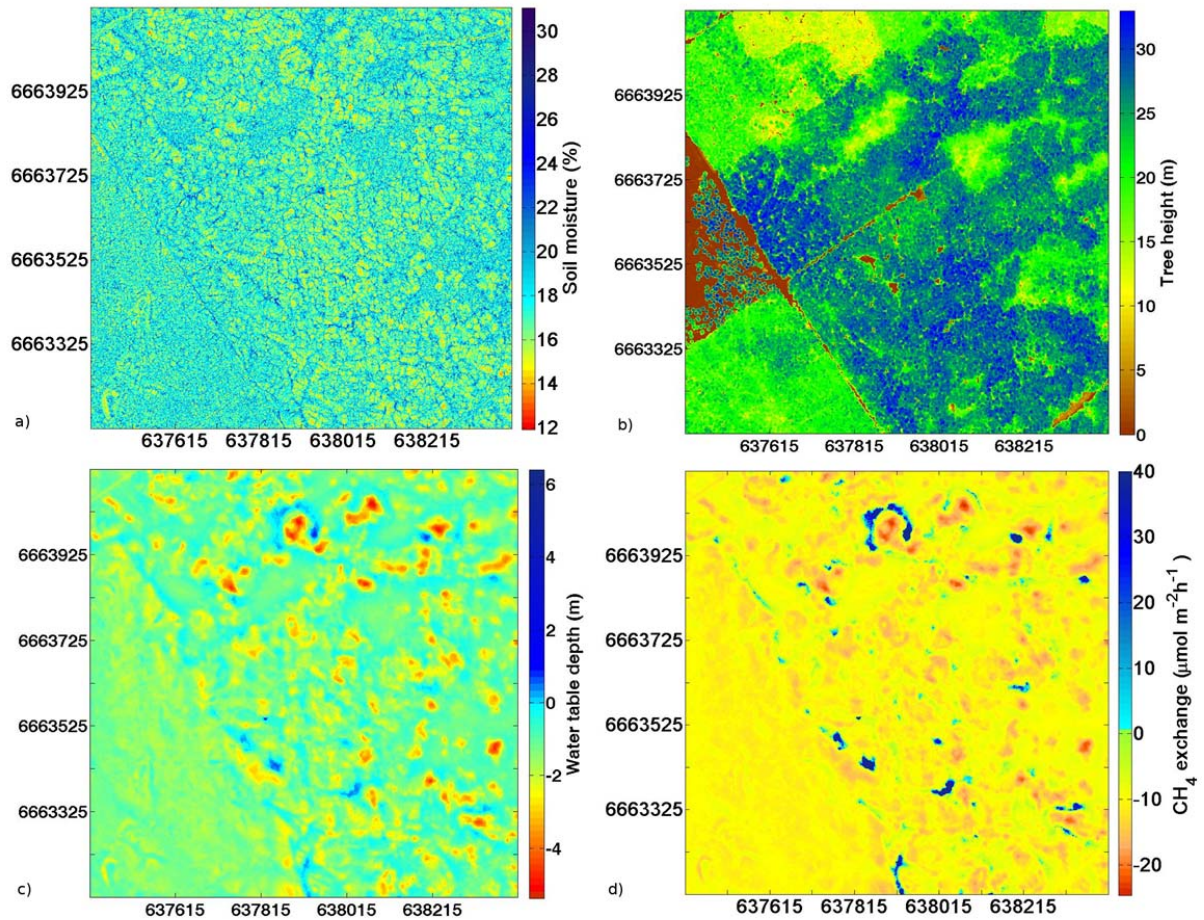


Figure 5. a) Map of soil moisture for the entire study area, averaged over the study time period July through September 2010. b) Map of tree height in the study area. c) Map of water table depth for the study area, averaged over the study time period. d) Map of modelled CH₄ exchange for the study area, averaged over the study time period. Coordinates are given in UTM (WGS 84).

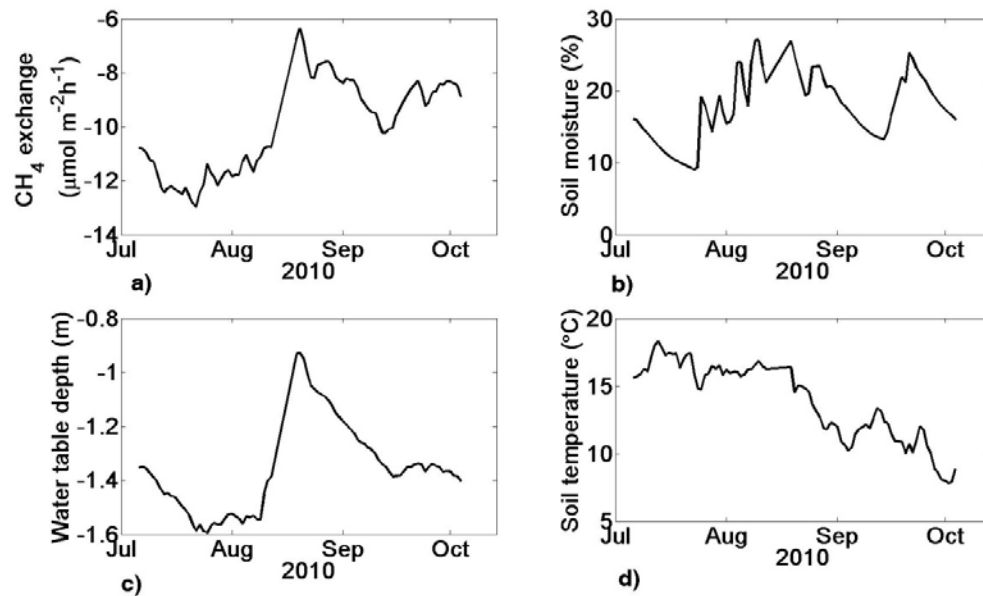


Figure 6. Time series of modelled a) CH₄ exchange, b) soil moisture, c) water table depth and d) soil temperature, spatially averaged over the study area.